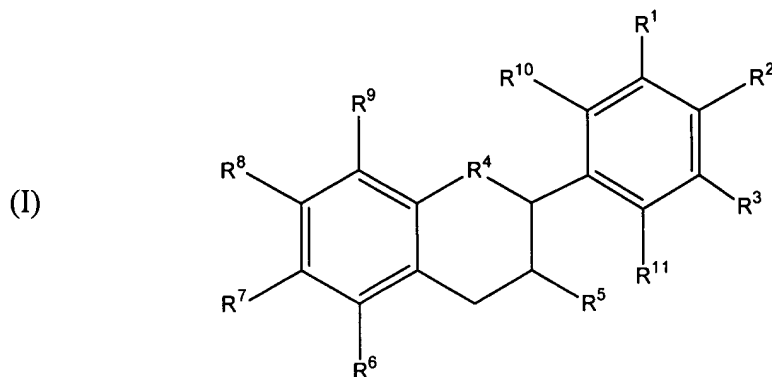


This listing of the claims replaces any and all prior versions and listings of claims in the application:

**LISTING OF THE CLAIMS**

1. (Currently amended) A compound having the structural formula (I)



wherein:

R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> are selected from the group consisting of ~~hydrogen~~, hydroxyl, alkyl, halo, sulfhydryl, alkoxy, and aryloxy, ~~and further wherein either R<sup>1</sup> and R<sup>2</sup>, or R<sup>2</sup> and R<sup>3</sup>, can be linked to form a cyclic group;~~

R<sup>4</sup> is selected from O, S, NR<sup>x</sup>, and CR<sup>y</sup>R<sup>z</sup>, wherein R<sup>x</sup>, R<sup>y</sup>, and R<sup>z</sup> are hydrogen or alkyl;

R<sup>5</sup> is selected from the group consisting of SH, acyloxy, and N(R<sup>x</sup>)<sub>2</sub> wherein the R<sup>x</sup> may be the same or different and are hydrogen or alkyl as defined previously;

R<sup>6</sup> and R<sup>8</sup> are independently selected from the group consisting of hydroxyl, alkyl, alkoxy, and aryloxy;

~~R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup> and R<sup>9</sup> are independently selected from the group consisting of hydrogen, hydroxyl, alkyl, alkoxy, and aryloxy, providing that either R<sup>6</sup> and R<sup>7</sup>, or R<sup>8</sup> and R<sup>9</sup>, may be linked together to form a cyclic structure selected from five-membered rings, six-membered rings, and fused five-membered and/or six-membered rings, wherein the cyclic structure is aromatic, alicyclic, heteroaromatic, or heteroalicyclic, and has zero to 4 non-hydrogen substituents and zero to 3 heteroatoms; and~~

$R^{10}$  and  $R^{11}$  are independently selected from the group consisting of hydrogen, hydroxyl, alkyl, alkoxy, and halo,

with the proviso that when (a)  $R^7$ ,  $R^9$ ,  $R^{10}$ , and  $R^{11}$  are hydrogen, and (b)  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^6$ , and  $R^8$  are hydroxyl, ~~(c)  $R^3$  is hydrogen or hydroxyl, and (d)  $R^4$  is O,~~ then ~~(e)(c)~~  $R^5$  is other than 3,4,5-trihydroxybenzoyloxy or 3,4,5-trimethoxybenzoyloxy.

2. (Currently amended) The compound of claim 1, wherein  $R^1$ ,  $R^2$  and  $R^3$  are selected from the group consisting of ~~hydrogen,~~ hydroxyl,  $C_1$ - $C_6$  alkyl, halo,  $C_1$ - $C_6$  alkoxy, and  $C_5$ - $C_{12}$  aryloxy, ~~and further wherein either  $R^1$  and  $R^2$ , or  $R^2$  and  $R^3$ , can be joined to form a two-atom or three-atom linkage selected from alkylene, substituted alkylene, and heteroalkylene;~~

~~$R^4$  is selected from O, S, NH and  $CH_2$ ;~~

$R^5$  is selected from the group consisting of  $C_6$ - $C_{32}$  acyloxy and  $NH_2$ ;

$R^6$  and  $R^8$  are independently selected from the group consisting of hydroxyl,  $C_1$ - $C_6$  alkyl,  $C_1$ - $C_6$  alkoxy, and  $C_5$ - $C_{12}$  aryloxy;

~~$R^6$ ,  $R^7$ ,  $R^8$  and  $R^9$  are independently selected from the group consisting of hydrogen, hydroxyl,  $C_1$ - $C_6$  alkyl,  $C_1$ - $C_6$  alkoxy, and  $C_5$ - $C_{12}$  aryloxy, or  $R^6$  and  $R^7$  are linked together to form a cyclohexyl, cyclopentyl, or phenyl ring, and  $R^8$  and  $R^9$  are hydrogen, or  $R^8$  and  $R^9$  are linked together to form a cyclohexyl, cyclopentyl, or phenyl ring, and  $R^6$  and  $R^7$  are hydrogen;~~  
and

$R^{10}$  and  $R^{11}$  are independently selected from the group consisting of hydrogen, hydroxyl,  $C_1$ - $C_6$  alkyl,  $C_1$ - $C_6$  alkoxy, and halo.

3. (Canceled).

4. (Currently amended) The compound of claim 2[[3]], wherein  $R^7$ ,  $R^9$ ,  $R^{10}$  and  $R^{11}$  are hydrogen.

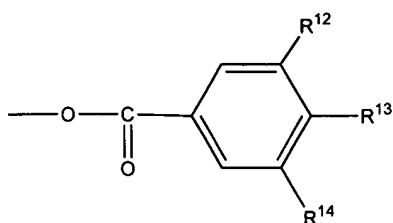
5. (Original) The compound of claim 4, in enantiomerically pure form in the 2 $\beta$ ,3 $\beta$ -*cis*, 2 $\alpha$ ,3 $\alpha$ -*cis*, 2 $\alpha$ ,3 $\beta$ -*trans*, or 2 $\beta$ ,3 $\alpha$ -*trans* configuration.

6. (Original) The compound of claim 4, comprising a racemic mixture of the 2 $\alpha$ ,3 $\beta$ -trans and 2 $\beta$ ,3 $\alpha$ -trans enantiomers.

7. (Original) The compound of claim 4, comprising a racemic mixture of the 2 $\alpha$ ,3 $\alpha$ -cis and 2 $\beta$ ,3 $\beta$ -cis enantiomers.

8. (Currently amended) The compound of claim 4, wherein:

R<sup>5</sup> is an acyloxy substituent having the structure



in which R<sup>12</sup>, R<sup>13</sup>, and R<sup>14</sup> are independently selected from the group consisting of hydrogen, hydroxyl, alkyl, alkoxy, and aryloxy, ~~such that the compound has the structure of formula (III).~~

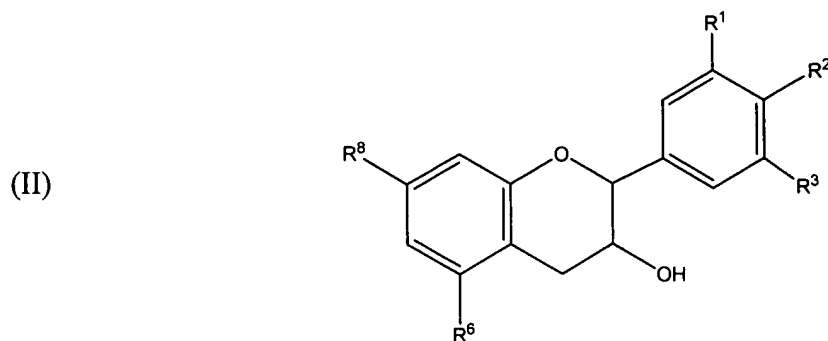
9. (Original) The compound of claim 8, wherein:

R<sup>12</sup>, R<sup>13</sup>, and R<sup>14</sup> are selected from the groups consisting of hydroxyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkoxy, and C<sub>5</sub>-C<sub>12</sub> aryloxy.

10. (Original) The compound of claim 9, wherein:

R<sup>12</sup>, R<sup>13</sup>, and R<sup>14</sup> are independently selected from the group consisting of hydroxyl, methyl, and methoxy, and benzyloxy.

11. (Currently amended) A compound having the structural formula (II)



wherein:

R<sup>1</sup>, R<sup>2</sup>, and R<sup>3</sup> are selected from the group consisting of hydroxyl, alkyl, halo, sulfhydryl, alkoxy, and aryloxy, ~~and further wherein either R<sup>1</sup> and R<sup>2</sup>, or R<sup>2</sup> and R<sup>3</sup>, can be linked to form a cyclic group;~~ and

R<sup>6</sup> and R<sup>8</sup> are selected from the group consisting of ~~hydrogen~~, alkyl, alkoxy, and aryloxy, wherein R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>6</sup> and R<sup>8</sup> are not all the same.

12. (Currently amended) The compound of claim 11, wherein R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> are selected from the group consisting of hydroxyl, C<sub>1</sub>-C<sub>6</sub> alkyl, halo, C<sub>1</sub>-C<sub>6</sub> alkoxy, and C<sub>5</sub>-C<sub>12</sub> aryloxy, ~~and further wherein either R<sup>1</sup> and R<sup>2</sup>, or R<sup>2</sup> and R<sup>3</sup>, can be joined to form a two atom or three atom linkage selected from alkylene, substituted alkylene, and heteroalkylene;~~ and

R<sup>6</sup> and R<sup>8</sup> are selected from the group consisting of ~~hydrogen~~, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkoxy, and C<sub>5</sub>-C<sub>12</sub> aryloxy.

13. (Currently amended) The compound of claim 12, wherein R<sup>1</sup>, R<sup>2</sup>, and R<sup>3</sup> are independently selected from hydroxyl, methyl, and methoxy; and

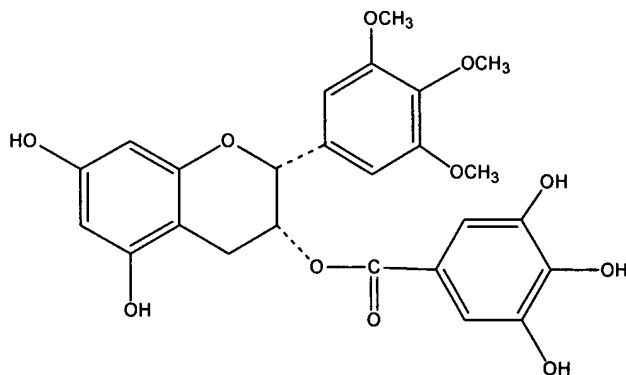
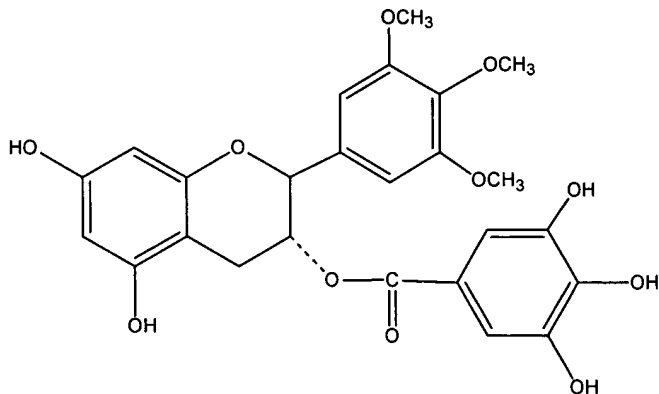
R<sup>6</sup>[[,]] and R<sup>8</sup> are independently selected from ~~hydrogen~~, hydroxyl, methyl, and methoxy.

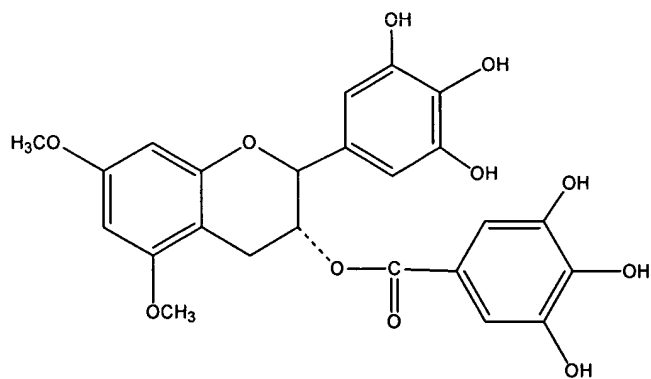
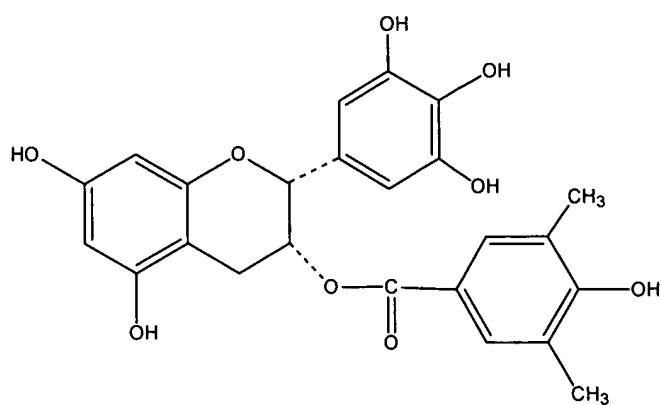
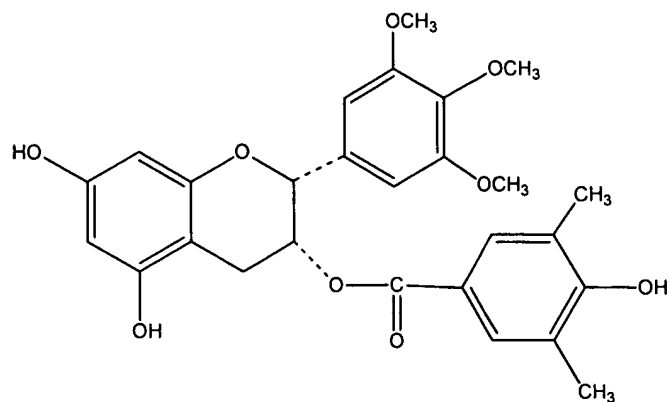
14. (Original) An analog of (-)-epigallocatechin-3-gallate (EGCG), wherein the analog contains at least one modification relative to ECGC that results in an IC<sub>50</sub> of less than 60 when the analog is evaluated for its ability to inhibit growth in a breast cancer cell line using MTT assay.

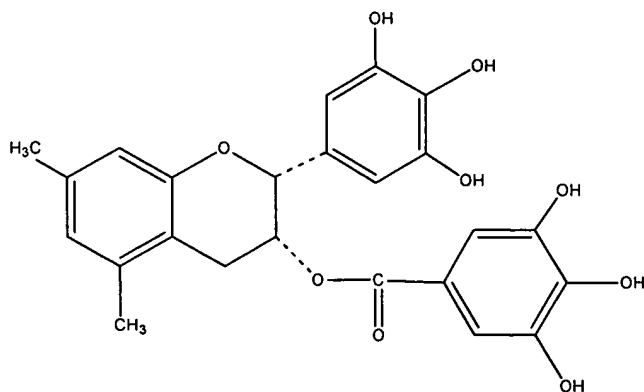
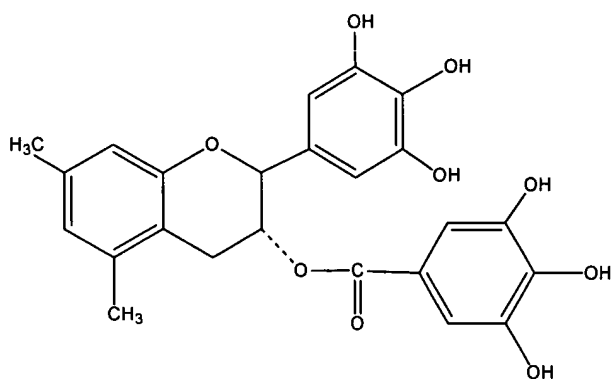
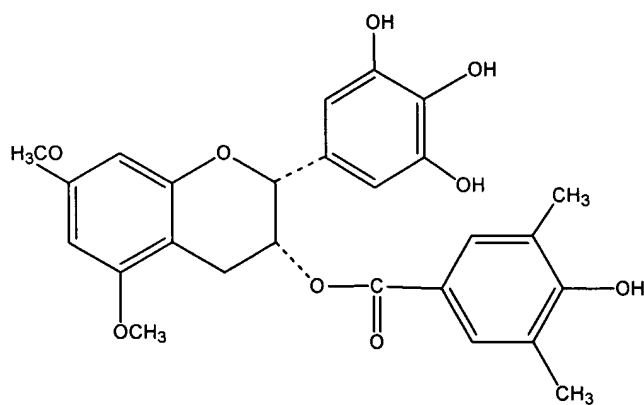
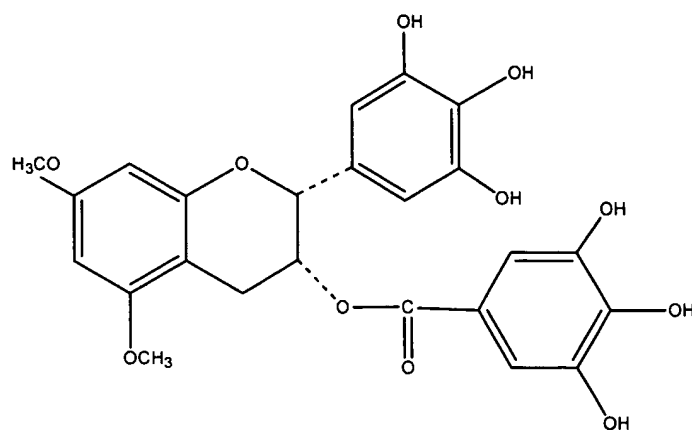
15. (Original) The analog of claim 14, wherein the analog contains at least one modification relative to ECGC that results in an  $IC_{50}$  of less than 25 when the analog is evaluated for its ability to inhibit growth in a breast cancer cell line using MTT assay.

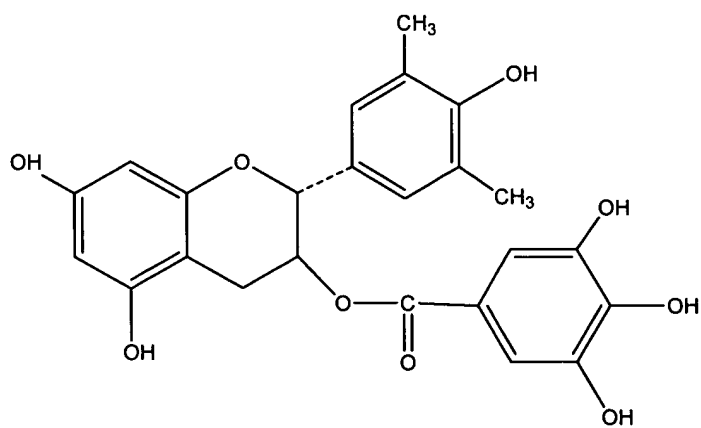
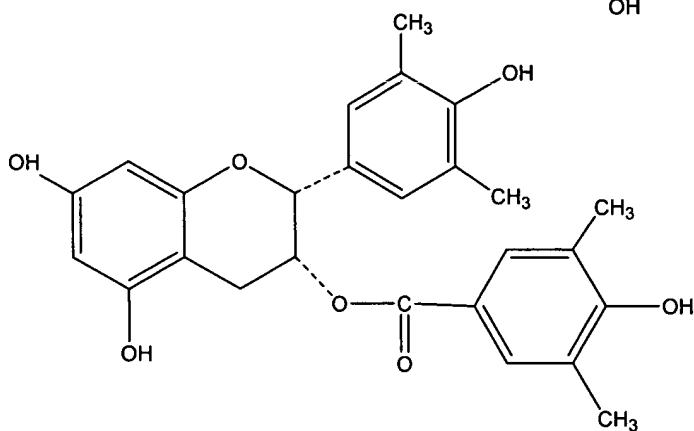
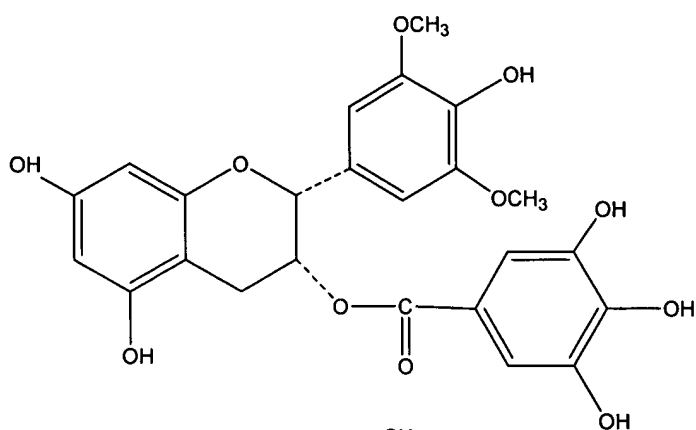
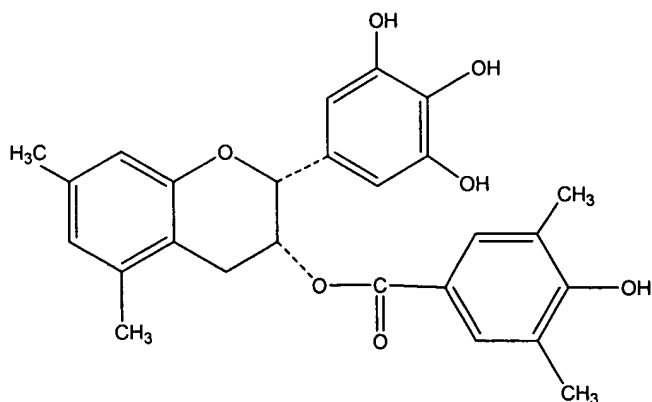
16. (Original) The analog of claim 15, wherein the analog contains at least one modification relative to ECGC that results in an  $IC_{50}$  of less than 15 when the analog is evaluated for its ability to inhibit growth in a breast cancer cell line using MTT assay.

17. (Original) A compound having the structural formula selected from

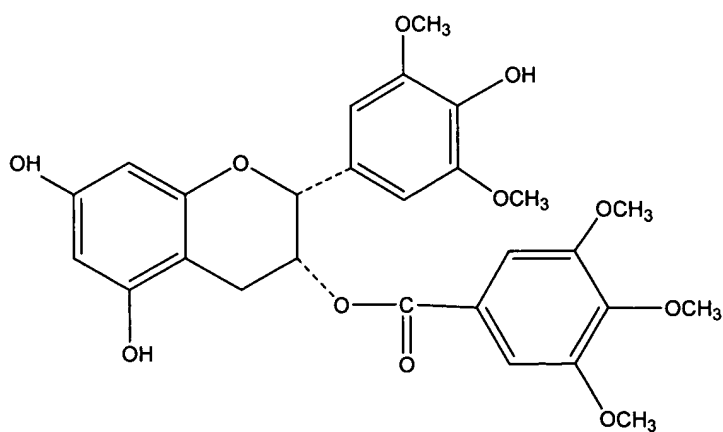
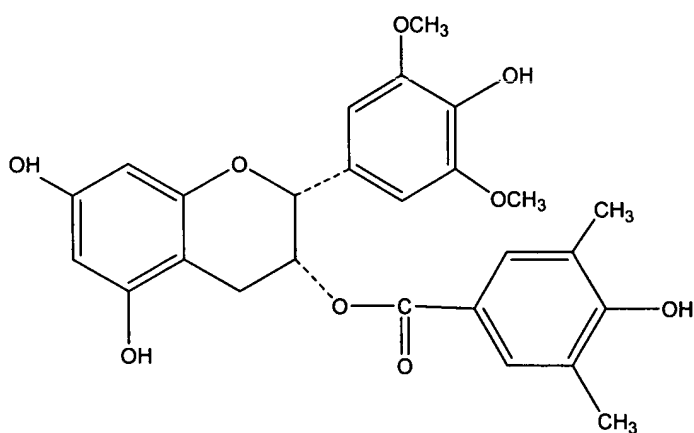
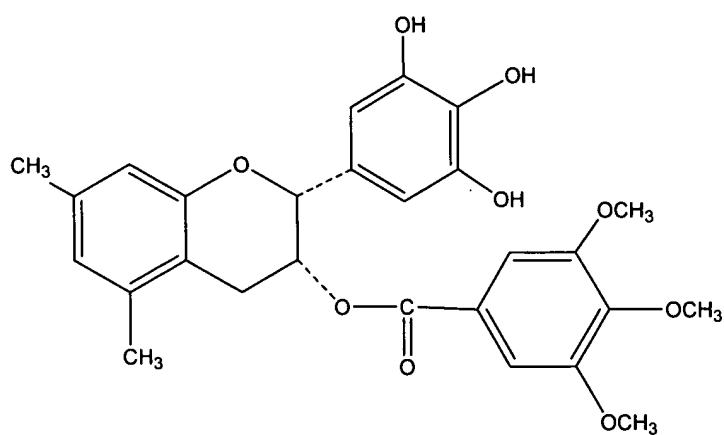
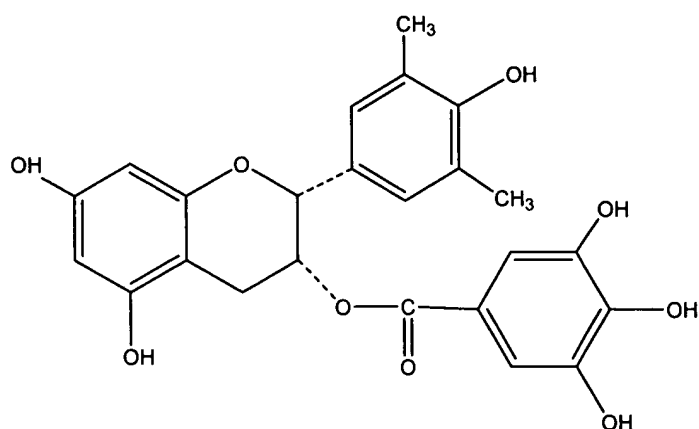












, and

18. (Original) A pharmaceutical composition comprising a therapeutically effective amount of the compound of claim 1 in combination with a pharmaceutically acceptable carrier.

19. (Currently amended) A pharmaceutical composition comprising a therapeutically effective amount of the compound of claim 2[[3]] in combination with a pharmaceutically acceptable carrier.

20. (Original) A pharmaceutical composition comprising a therapeutically effective amount of the compound of claim 4 in combination with a pharmaceutically acceptable carrier.

21. (Original) A pharmaceutical composition comprising a therapeutically effective amount of the compound of claim 5 in combination with a pharmaceutically acceptable carrier.

22. (Original) A pharmaceutical composition comprising a therapeutically effective amount of the compound of claim 6 in combination with a pharmaceutically acceptable carrier.

23. (Original) A pharmaceutical composition comprising a therapeutically effective amount of the compound of claim 8 in combination with a pharmaceutically acceptable carrier.

24. (Original) A pharmaceutical composition comprising a therapeutically effective amount of the compound of claim 9 in combination with a pharmaceutically acceptable carrier.

25. (Original) A pharmaceutical composition comprising a therapeutically effective amount of the compound of claim 15 in combination with a pharmaceutically acceptable carrier.

26. (Original) The composition of any one of claims 18 through 25, wherein the pharmaceutically acceptable carrier is suitable for oral administration and the composition comprises an oral dosage form.

27. (Original) The composition of claim 26, wherein the oral dosage form is a tablet.
28. (Original) The composition of claim 26, wherein the oral dosage form is a capsule.
29. (Original) The composition of any one of claims 18 through 25, wherein the pharmaceutically acceptable carrier is suitable for parenteral administration and the composition comprises a parenterally administrable formulation.
30. (Withdrawn) A method for treating a patient suffering from cancer, comprising administering to the individual a therapeutically effective amount of the compound of claim 1.
31. (Withdrawn) A method for treating a patient suffering from cancer, comprising administering to the individual a therapeutically effective amount of the compound of claim 3.
32. (Withdrawn) A method for treating a patient suffering from cancer, comprising administering to the individual a therapeutically effective amount of the compound of claim 4.
33. (Withdrawn) A method for treating a patient suffering from cancer, comprising administering to the individual a therapeutically effective amount of the compound of claim 5.
34. (Withdrawn) A method for treating a patient suffering from cancer, comprising administering to the individual a therapeutically effective amount of the compound of claim 6.
35. (Withdrawn) A method for treating a patient suffering from cancer, comprising administering to the individual a therapeutically effective amount of the compound of claim 8.
36. (Withdrawn) A method for treating a patient suffering from cancer, comprising administering to the individual a therapeutically effective amount of the compound of claim 9.
37. (Withdrawn) A method for treating a patient suffering from cancer, comprising administering to the individual a therapeutically effective amount of the compound of claim 15.

38. (Withdrawn) The method of any one of claims 30 through 37, wherein the cancer is prostate cancer, uterine cancer, or breast cancer.

39. (Withdrawn) The method of claim 38, wherein the cancer is breast cancer.

40. (Withdrawn) A chemopreventive method comprising administering a prophylactically effective amount of the compound of claim 1 to a patient susceptible to developing cancer.

41. (Withdrawn) A chemopreventive method comprising administering a prophylactically effective amount of the compound of claim 3 to a patient susceptible to developing cancer.

42. (Withdrawn) A chemopreventive method comprising administering a prophylactically effective amount of the compound of claim 4 to a patient susceptible to developing cancer.

43. (Withdrawn) A chemopreventive method comprising administering a prophylactically effective amount of the compound of claim 5 to a patient susceptible to developing cancer.

44. (Withdrawn) A chemopreventive method comprising administering a prophylactically effective amount of the compound of claim 6 to a patient susceptible to developing cancer.

45. (Withdrawn) A chemopreventive method comprising administering a prophylactically effective amount of the compound of claim 8 to a patient susceptible to developing cancer.

46. (Withdrawn) A chemopreventive method comprising administering a prophylactically effective amount of the compound of claim 9 to a patient susceptible to developing cancer.

47. (Withdrawn) A chemopreventive method comprising administering a prophylactically effective amount of the compound of claim 15 to a patient susceptible to developing cancer.